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Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1-16. (Canceled)

17. (Currently Amended) A medicament method for preventive and/or therapeutic treatment of dermal pigmentation and/or development of skin cancer in a mammal including a human, which comprises the step of administering a preventively and/or therapeutically effective amount of as an active ingredient a substance selected from the group consisting of a compound represented by the following general formula (I) and a pharmacologically acceptable salt thereof, and a hydrate thereof and a solvate thereof:

wherein X represents a group represented by the following formula:

wherein a bond at the left end binds to ring Z and a bond at the right end binds to E,

A represents hydrogen atom,

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E represents a 2,5-di-substituted phenyl group wherein at least one of said substituents is trifluoromethyl group, a 3,5-di-substituted phenyl group wherein at least one of said substituents is trifluoromethyl group, or a 4,5-di-substituted thiazol-2-yl group,

ring Z represents a C₆ to C₁₀ arene which may have one or more substituents in addition to the group represented by formula –O-A wherein A has the same meaning as that defined above and the group represented by formula –X-E wherein each of X and E has the same meaning as that defined above, or a 5- to 13-membered heteroarene which may have one or more substituents in addition to the group represented by formula –O-A wherein A has the same meaning as that defined above and the group represented by formula –X-E wherein each of X and E has the same meaning as that defined above, to a mammal including a human.

18. (Currently Amended): The medicament method according to claim 17, wherein E is a group selected from the group consisting of the following substituent group δ -3e, substituent group δ -5e, and substituent group δ -8e, the following partial formula (Iz-1) in the general formula (I) containing ring Z

is the following formula (Iz-2):

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wherein R^z represents a group selected from the following substituent group γ -2z[[.]], wherein

FSubstituent Group δ-3e[†] represents a 2-chloro-5-(trifluoromethyl)phenyl group, a 2,5-

bis(trifluoromethyl)phenyl group, a 2-fluoro-5-(trifluoromethyl)phenyl group, a 2-nitro-5-

(trifluoromethyl)phenyl group, a 2-methyl-5-(trifluoromethyl)phenyl group, a 2-methoxy-5-

(trifluoromethyl)phenyl group, a 2-methylsulfanyl-5-(trifluoromethyl)phenyl group, a 2-(1-

pyrrolidinyl)-5-(trifluoromethyl)phenyl group, a 2-morpholino-5-(trifluoromethyl)phenyl group, a

2-bromo-5-(trifluoromethyl)phenyl group, a 2-(2-naphthyloxy)-5-(trifluoromethyl)phenyl group,

a 2-(2,4-dichlorophenoxy)-5-(trifluoromethyl)phenyl group, a 2-[4-(trifluoromethyl)piperidin-1-

yll-5-(trifluoromethyl)phenyl group, a 2-(2,2,2-trifluoroethoxy)-5-(trifluoromethyl)phenyl group,

a 2-(2-methoxyphenoxy)-5-(trifluoromethyl)phenyl group, a 2-(4-chloro-3,5-dimethylphenoxy)-

5-(trifluoromethyl)phenyl group, a 2-piperidino-5-(trifluoromethyl)phenyl group, a 2-(4-

methylphenoxy)-5-(trifluoromethyl)phenyl group, a 2-(4-chlorophenoxy)-5-

(trifluoromethyl)phenyl group, a 2-(4-cyanophenoxy)-5-(trifluoromethyl)phenyl group, a 2-(4-

methoxyphenoxy)-5-(trifluoromethyl)phenyl group;

fSubstituent Group δ-5el represents a 3,5-bis(trifluoromethyl)phenyl group, a 3-fluoro-5-

(trifluoromethyl)phenyl group, a 3-bromo-5-(trifluoromethyl)phenyl group, a 3-methoxy-5-

(trifluoromethyl)phenyl group, a 3-methoxycarbonyl-5-(trifluoromethyl)phenyl group, a 3-

carboxy-5-(trifluoromethyl)phenyl group;

FSubstituent Group δ-8e] represents a 5-bromo-4-[(1,1-dimethyl)ethyl]thiazol-2-yl group, a 5-

bromo-4-(trifluoromethyl)thiazol-2-yl group, a 5-cyano-4-[(1,1-dimethyl)ethyl]thiazol-2-yl

group, a 4,5-dimethylthiazol-2-yl group, a 5-methyl-4-phenylthiazol-2-yl group, a 5-(4-

fluorophenyl)-4-methylthiazol-2-yl group, a 4-methyl-5-[3-(trifluoromethyl)phenyl]thiazol-2-yl

group, a 4-[(1,1-dimethyl)ethyl]-5-ethylthiazol-2-yl group, a 4-ethyl-5-phenylthiazol-2-yl group,

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<u>a</u> 4-isopropyl-5-phenylthiazol-2-yl group, <u>a</u> 4-butyl-5-phenylthiazol-2-yl group, <u>a</u> 4-[(1,1dimethyl)ethyl]-5-[(2,2-dimethyl)propionyl]thiazol-2-yl group, a 4-[(1,1-dimethyl)ethyl]-5-(ethoxycarbonyl)thiazol-2-yl group, a 4-[(1,1-dimethyl)ethyl]-5-piperidinothiazol-2-yl group, a 4-[(1,1-dimethyl)ethyl]-5-morpholinothiazol-2-yl group, a 4-[(1,1-dimethyl)ethyl]-5-(4methylpiperazin-1-yl)thiazol-2-yl group, a 4-[(1,1-dimethyl)ethyl]-5-(4-phenylpiperazin-1yl)thiazol-2-yl group, a 5-carboxymethyl-4-phenylthiazol-2-yl group, a 4,5-diphenylthiazol-2-yl group, 4-benzyl-5-phenylthiazol-2-yl group, a 5-phenyl-4-(trifluoromethyl)thiazol-2-yl group, a 5-acetyl-4-phenylthiazol-2-yl group, a 5-benzoyl-4-phenylthiazol-2-yl group, a 5-ethoxycarbonyl-4-phenylthiazol-2-yl group, a 5-ethoxycarbonyl-4-(pentafluorophenyl)thiazol-2-yl group, a 5methylcarbamoyl-4-phenylthiazol-2-yl group, a 5-ethylcarbamoyl-4-phenylthiazol-2-yl group, a 5-isopropylcarbamoyl-4-phenylthiazol-2-yl group, a 5-(2-phenylethyl)carbamoyl-4phenylthiazol-2-yl group, a 5-ethoxycarbonyl-4-(trifluoromethyl)thiazol-2-yl group, a 5-carboxy-4-[(1,1-dimethyl)ethyl]thiazol-2-yl group, a 5-(ethoxycarbonyl)methyl-4-phenylthiazol-2-yl group, a 5-carboxy-4-phenylthiazol-2-yl group, a 5-propylcarbamoyl-4-phenylthiazol-2-yl group; {Substituent Group γ -2z} represents a halogen atom, a nitro group, a cyano group, a methoxy group, a methyl group, an isopropyl group, a tert-butyl group, a 1,1,3,3-tetramethylbutyl group, a 2-phenylethen-1-yl group, a 2,2-dicyanoethen-1-yl group, a 2-cyano-2-(methoxycarbonyl)ethen-1-yl group, <u>a</u> 2-carboxy-2-cyanoethen-1-yl group, <u>an</u> ethynyl group, <u>a</u> phenylethynyl group, (trimethylsilyl)ethynyl group, <u>a</u> trifluoromethyl group, <u>a</u> pentafluoroethyl group, <u>a</u> phenyl group, a 4-(trifluoromethyl)phenyl group, a 4-fluorophenyl group, a 2,4-difluorophenyl group, a 2phenethyl group, a 1-hydroxyethyl group, a 1-(methoxyimino)ethyl group, a 1-[(benzyloxy)imino]ethyl group, a 2-thienyl group, a 3-thienyl group, a 1-pyrrolyl group, a 2methylthiazol-4-yl group, an imidazo[1,2-a]pyridin-2-yl group, a 2-pyridyl group, an acetyl

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group, <u>an</u> isobutyryl group, <u>a</u> piperidinocarbonyl group, <u>a</u> 4-benzylpiperidinocarbonyl group, <u>a</u> (pyrrol-1-yl)sulfonyl group, <u>a</u> carboxy group, methoxycarbonyl group, <u>an</u> N-[3,5-bis(trifluoromethyl)phenyl]carbamoyl group, <u>an</u> N,N-dimethylcarbamoyl group, <u>a</u> sulfamoyl group, an N-[3,5-bis(trifluoromethyl)phenyl]sulfamoyl group, an N,N-dimethylsulfamoyl group,

amino group, an N,N-dimethylamino group, an acetylamino group, a benzoylamino group, a

methanesulfonylamino group, <u>a</u> benzenesulfonylamino group, <u>a</u> 3-phenylureido group, <u>a</u> (3-

phenyl)thioureido group, a (4-nitrophenyl)diazenyl group, a {[4-(pyridin-2-

yl)sulfamoyl]phenyl}diazenyl group.

19. (Currently Amended) The medicament method according to claim 18, wherein E is a group selected from the group consisting of the aforementioned substituent group δ -3e, substituent group δ -5e, and substituent group δ -8e, and R^z is a halogen atom.

20. (Currently Amended) The medicament method according to claim 19, wherein E is 2,5-bis(trifluoromethyl)phenyl group or 3,5-bis(trifluoromethyl)phenyl group, and R^z is a halogen atom.

- 21. (Currently Amended) The medicament method according to claim 20, wherein E is 3,5-bis(trifluoromethyl)phenyl group, and R^z is a halogen atom.
- 22. (Currently Amended): The medicament method according to claim 17, wherein E is 2,5-bis(trifluoromethyl)phenyl group or 3,5-bis(trifluoromethyl)phenyl group.

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23. (Currently Amended) The medicament method according to claim 17,

wherein the following partial formula (Iz-1) in the general formula (I) containing ring Z

is the following formula (Iz-2):

wherein R^z represents a halogen atom.

- 24. (Currently Amended) The medicament method according to claim 17, for preventive and/or therapeutic treatment of dermal pigmentation.
- 25. (Currently Amended) The medicament method according to claim 21, for preventive and/or therapeutic treatment of dermal pigmentation.
- 26. (Currently Amended) The medicament according to claim 17, having inhibitory activity against A method of inhibiting transformation and/or proliferation of melanocytes caused by ultraviolet irradiation, which comprises the step of allowing a substance according to claim 17 to act on melanocytes.

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27. (Currently Amended) The medicament method according to claim 26, wherein the substance is the same as that defined in claim 21, having inhibitory activity against transformation and/or proliferation of melanocytes caused by ultraviolet irradiation.

- 28. (Currently Amended) A cosmetic composition which comprises a substance according to claim 17 as an active ingredient having a method of skin whitening effect, which comprises the step of administering an effective amount of a substance according to claim 17 to a human.
- 29. (Currently Amended) A cosmetic composition which comprises a substance The method according to claim 28, wherein the substance is the same as that defined in claim 21 as an active ingredient having a skin whitening effect.
 - 30. (New) The method according to claim 17, wherein the mammal is a human.